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**FACSIMILE COVER SHEET****Licata & Tyrrell P.C.**

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October 21, 2003

**TO: Examiner Horlick (TC1600)****GROUP: 1637****FAX NUMBER: 703-872-9306****ATTORNEY DOCKET NO.: DEX-0269****SERIAL NO.: 10/001,887****FILED: November 20, 2001****NUMBER OF PAGES:**

**MESSAGE:** Attached please find Amendment Transmittal Letter, Reply to Restriction Requirement and Certificate of Transmission by Facsimile.

**Kathleen A. Tyrrell, Registration No. 38,350****URGENT! PLEASE DELIVER IMMEDIATELY UPON RECEIPT. THANK YOU!**

\* \* \* \* \*

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**CERTIFICATE OF TRANSMISSION BY FACSIMILE (37 CFR 1.8)**

Applicant(s): Salceda et al.

Docket No.

DEX-0269

Serial No.

10/001,887

Filing Date

November 20, 2001

Examiner

Horlick, Kenneth R.

Group Art Unit

1637

Invention:

**Compositions and Methods Relating to Breast Specific Genes and Proteins**

I hereby certify that this Reply to Restriction Requirement

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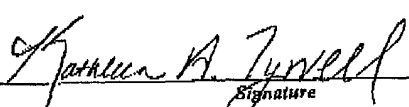
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AMENDMENT TRANSMITTAL LETTER (Large Entity)				Docket No. DEX-0269	
Applicant(s): Salceda et al.					
Serial No. 10/001,887	Filing Date November 20, 2001	Examiner Horlick, Kenneth R.		Group Art Unit 1637	
Invention: Compositions and Methods Relating to Breast Specific Genes and Proteins					
<u>TO THE COMMISSIONER FOR PATENTS:</u>					
Transmitted herewith is an amendment in the above-identified application. The fee has been calculated and is transmitted as shown below.					
CLAIMS AS AMENDED					
	CLAIMS REMAINING AFTER AMENDMENT	HIGHEST # PREV. PAID FOR	NUMBER EXTRA CLAIMS PRESENT	RATE	ADDITIONAL FEE
TOTAL CLAIMS	19 -	20 =	0 x	\$18.00	\$0.00
INDEP. CLAIMS	2 -	3 =	0 x	\$86.00	\$0.00
Multiple Dependent Claims (check if applicable) <input type="checkbox"/>					\$0.00
TOTAL ADDITIONAL FEE FOR THIS AMENDMENT					\$0.00
<input checked="" type="checkbox"/> No additional fee is required for amendment. <input type="checkbox"/> Please charge Deposit Account No. _____ in the amount of _____ <input type="checkbox"/> A check in the amount of _____ to cover the filing fee is enclosed. <input checked="" type="checkbox"/> The Director is hereby authorized to charge payment of the following fees associated with this communication or credit any overpayment to Deposit Account No. 50-1619 <input checked="" type="checkbox"/> Any additional filing fees required under 37 C.F.R. 1.16. <input checked="" type="checkbox"/> Any patent application processing fees under 37 CFR 1.17.					
 Kathleen A. Tyrrell, Reg. No. 38,350			Dated: October 21, 2003		
Licata & Tyrrell P.C. 66 East Main Street Marlton, New Jersey 08053 Tel : 856-810-1515 Fax: 856-810-1454			<div>I certify that this document and fee is being deposited on _____ with the U.S. Postal Service as first class mail under 37 C.F.R. 1.8 and is addressed to the Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.</div> <div>Signature of Person Mailing Correspondence</div> <div>Typed or Printed Name of Person Mailing Correspondence</div>		
cc:					

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Attorney Docket No.: DEX-0269  
Inventors: Salceda et al.  
Serial No.: 10/001,887  
Filing Date: November 20, 2001  
Examiner: Horlick, Kenneth R.  
Group Art Unit: 1637  
Title: Compositions and Methods Relating to  
Breast Specific Genes and Proteins

## Certificate of Facsimile Transmission

I hereby certify that this document is being facsimile  
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the date shown below.

On October 21, 2003

  
Kathleen A. Tyrrell, Registration No. 38,350

Mail Stop  
Commissioner for Patents  
PO Box 1450  
Alexandria, VA 22313-1450

Dear Sir:

Reply to Restriction Requirement

This is a reply to the Restriction Requirement mailed  
September 22, 2002 setting a one (1) month statutory period for  
response. Please enter the following remarks into the record.

Remarks begin on page 2.

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Attorney Docket No.: DEX-0269  
Inventors: Salceda et al.  
Serial No.: 10/001,887  
Filing Date: November 20, 2001  
Page 2

#### REMARKS

Claims 1-17 are pending in the instant application. Claims 1-17 have been subjected to a Restriction Requirement as follows:

Group I, claims 1-5, 7-9 and 15 (partial), drawn to nucleic acids, vectors, host cells, and methods of making a polypeptide, classified in class 536, subclass 23.1, and class 435, subclasses 69.1, 320.1 and 325, for example;

Group II, claim 10-11, drawn to polypeptides, classified in class 530, subclass 350, for example;

Group III, claims 12 and 15 (partial), drawn to an antibody, classified in class 530, subclass 387.1, for example;

Group IV, claims 6 and 14 (partial), drawn to a method for determining the presence of a nucleic acid, classified in class 435, subclass 6;

Group V, claims 13 and 14 (partial), drawn to a method for determining the presence of a polypeptide, classified in class 435, subclass 7.1, for example;

Group VI, claim 16, drawn to a method for treating a patient with breast cancer by administering an antibody, classified in class 514, subclass 2, for example;

Group VII, claim 17(partial), drawn to a vaccine comprising

Attorney Docket No.: DEX-0269  
Inventors: Salceda et al.  
Serial No.: 10/001,887  
Filing Date: November 20, 2001  
Page 3

a polypeptide, classified in class 514, subclass 2; and  
Group VIII, claim 17 (partial), drawn to a vaccine  
comprising a nucleic acid, classified in class 514, subclass 44,  
for example.

The Examiner suggests that these Groups are distinct.

Specifically, with respect to Groups I, II, III, VII and  
VIII, the Examiner suggests that the claims are drawn to  
different products having different structures and functions.

With respect to Groups I and IV, and Groups III and (V,VI),  
the Examiner has acknowledged their relationship as product and  
process of use. However, the Examiner suggests that the Groups  
are distinct because the products can be used in materially  
different methods or processes.

With respect to Groups I and (V, VI), Groups II and (IV, V  
and VI), Groups III and IV, Groups IV-VI, and Groups (IV-VI) and  
(VII,VIII), the Examiner suggests that the Groups are unrelated  
because the different Groups are not required for one another.

Further, the Examiner suggests that each of Groups I-VIII  
are drawn to a multitude of nucleic acids, polypeptides, and  
antibodies thereto which are independent and distinct. Thus, the  
Examiner is also requiring election of a single nucleic acid,  
polypeptide or antibody.

Attorney Docket No.: DEX-0269  
Inventors: Salceda et al.  
Serial No.: 10/001,887  
Filing Date: November 20, 2001  
Page 4

Applicants respectfully traverse this Restriction Requirement.

MPEP §803 provides two criteria which must be met for a restriction requirement to be proper. The first is that the inventions be independent or distinct. The second is that there would be a serious burden on the Examiner if the restriction is not required. A search of prior art relating to an elected nucleic acid, polypeptide or antibody would also reveal any references teaching uses for the nucleic acid, polypeptide or antibody. Accordingly, Applicants believe that searching of all the claims, at least when limited to elected nucleic acids or polypeptides is overlapping and would not place an undue burden on the Examiner if the Restriction is not made.

Thus, since this Restriction Requirement does not meet both criteria as set forth in MPEP § 803 to be proper, reconsideration and withdrawal of this Restriction Requirement is respectfully requested.

In addition, with respect to the election of a single sequence, MPEP § 803.04 clearly states that a reasonable number of nucleotide sequences, normally ten sequences, can be claimed in a single application. Accordingly, withdrawal of this sequence election requirement and reconsideration to include a

Attorney Docket No.: DEX-0269  
Inventors: Salceda et al.  
Serial No.: 10/001,887  
Filing Date: November 20, 2001  
Page 5

more reasonable number of at least 10 sequences in accordance with MPEP § 803.04 is also respectfully requested.

However, in an earnest effort to advance the prosecution of this case, Applicants elect Group I, claims 1-5, 7-9 and 15, SEQ ID NO:64 encoding SEQ ID NO:127, with traverse.

Applicants believe that the foregoing comprises a full and complete response to the Office Action of record.

Respectfully submitted,

*Kathleen A. Tyrrell*  
Kathleen A. Tyrrell  
Reg. No. 38,350

Date: October 21, 2003

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